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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C07K 7/06, 7/08, 7/10 G01N 33/569		A1	(11) International Publication Number: WO 93/14118 (43) International Publication Date: 22 July 1993 (22.07.93)
(21) International Application Number: PCT/GB93/00087 (22) International Filing Date: 15 January 1993 (15.01.93) (30) Priority data: 9200949.7 17 January 1992 (17.01.92) GB (71) Applicant (for all designated States except US): MEDICAL RESEARCH COUNCIL [GB/GB]; 20 Park Crescent, London W1N 4AL (GB). (72) Inventors; and (75) Inventors/Applicants (for US only) : IVANYI, Juraj [GB/GB]; ELSAGHIER, Ashraf [EG/GB]; MRC Tuberculosis and Related Infections Unit, Royal Postgraduate Medical School, Hammersmith Hospital, Du Cane Road, London W12 0HS (GB).		(74) Agents: CRESSWELL, Thomas, Anthony et al.; J.A. Kemp & Co., 14 South Square, Gray's Inn, London WC1R 5EU (GB). (81) Designated States: AU, CA, JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	

(54) Title: PEPTIDE FRAGMENTS OF HSP71 OF *M. TUBERCULOSIS* AND THEIR USE IN DIAGNOSIS OF TUBERCULOSIS

Hu: A K . A L . S Y A F N M . S A . . D E G L K
Ec: T . . G D H . L H S . R . Q . E . A G D K
Lp: T
Tb: V R N Q A E I L V Y Q I E K E V K E Q R E A 530

- . - - - I S - A D K K - . L D K C Q - V I S W
 L P A D D K T A I E S A L T A L E T - L K G E -
 - N - R - E - E T A
 E G G S K V P E D I T L N K V D A A V A E A E G - 554

L D A N T L - A E K D E F E H K R K E L E L Q V
- D K A A . E A K M Q E L A Q V S Q K L M E I .
L . G T D . - S A I K S A M E K L G Q D S Q A L
- G T W R I - G Y F G H Q V G D G E A G P G V A 578

C N P I I S G L Y Q G A G G - - P G P G G F G A
Q Q O H A Q Q Q T A G A D A S A N N A K D - - -
• Q A I Y E A T Q A A S K • G • E A S A • G G S
G S G A S D L R S S S G C V I G H W R C P - - - 602

Q G P K G - G S G S G P T I E E V D -
 - - - D D V V D A E F E E V K D - K K
 N S T D D V L T . . W S T T N G S P K
 - - - - - P R R R A G R C P P R L G 621

(57) Abstract

Peptides comprising a sequence the same as or immunologically equivalent to a linear epitope of the carboxy terminal region of *Mycobacteria tuberculosis* heat shock proteins such as Hsp71 useful in the diagnosis of paucibacillary tuberculosis (TB). Methods of diagnosis of TB using such peptides and test kits including such peptides are also provided.

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PEPTIDE FRAGMENTS OF HSP71 OF M. TUBERCULOSIS AND THEIR USE IN DIAGNOSIS OF TUBERCULOSIS

The present invention relates to peptides which are useful in diagnosing paucibacillary mycobacterial infections.

5 The paucibacillary stage of tuberculosis is very hard to detect or diagnose due to the effective absence of clinical signs and symptoms and the lack of detectable pathogen-originating antigens. Diagnosis in the multibacillary stage is relatively easy, for instance using
10 well known smear test. Diagnosis at the paucibacillary (or smear-negative) stage is important in order that appropriate treatment and containment measures can be implemented to avoid transmission to persons at the greatest risk infection. To date efforts to improve the
15 diagnosis of paucibacillary TB have been unsuccessful.

 The present inventors have now identified certain peptides which may be used in detecting paucibacillary TB and other mycobacterial diseases. The peptides are related to the carboxy terminal of the stress
20 protein Hsp71 of Mycobacterium tuberculosis and corresponding heat shock proteins of other mycobacterial pathogens.

Without wishing to be bound by this theory the

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present inventors consider that the carboxy terminal regions of Hsp71 and corresponding mycobacterial heat shock proteins contain species-specific determinants having linear epitopes. These linear epitopes generate an antibody response during the paucibacillary stage of infection and peptides having corresponding sequences can be used to detect the antibodies in the serum or other body fluids of the patient.

Peptides of the invention may be used for detecting or diagnosing a variety of paucibacillary mycobacterial diseases especially, but not limited to, tuberculosis of humans and wild or domestic mammals such as cattle, donkeys, deer and badgers. The peptides may also be used in detecting or diagnosing multibacillary mycobacterial infections.

The present invention therefore provides a peptide comprising a sequence the same as or immunologically equivalent to a linear epitope of the carboxy terminal region of M. tuberculosis Hsp71 or a corresponding heat shock protein of any other mycobacterial pathogen.

Preferably the peptide comprises a sequence the same as or immunologically equivalent to a linear epitope of the carboxy terminal of M. tuberculosis Hsp71.

The invention will be illustrated with reference to the figures of the accompanying drawings in

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which:-

5 **Figure 1** shows the aligned carboxy terminal region
sequences of M. tuberculosis Hsp71 and
corresponding heat shock proteins using the
internationally-accepted 1-letter code for
amino acid residues. Hu=Human hsp70-1,
Ec=Escherichia coli dnaK, Lp=Mycobacterium
10 leprae Hsp71, Tb=Mycobacterium tuberculosis
Hsp71 .=sequences identical with
M.tuberculosis. -=gaps.

Figure 2 shows an analysis of antigens by SDS PAGE and
Western blots. Ec=extract from E coli
a=coomassie blue stain; b=Western blots
developed with tuberculosis patient's serum.

15 **Figure 3** gives individual serum antibody titers to
mycobacterial hsp70 (M-hsp70) and human
hsp70(H-hsp70 in groups of patients and
controls. Groups:active TB(A); "self-healed"
inactive TB(B); past treated inactive TB (C);
20 lung fibrosis (D); diseases other than
tuberculosis (E); healthy BCG-vaccinated
individuals (F) (see also table 1). Group means
are indicated by the horizontal bars. Boxed
symbols in group B: Patients given

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chemoprophylaxis due to severity of chest X-ray and clinical evaluation.

Figure 4 gives antibody levels in active tuberculosis divided according to sputum bacillary status.

5 S=smear microscopy; C=culture of M tuberculosis. Group means are indicated by horizontal bars.

Figure 5 gives the ratios between anti-mycobacterial and anti-human hsp70 antibody levels. Groups of patients:see legend to table 1.

10

Figure 6 Shows antibody binding to octamer peptides derived from the carboxy-terminal sequence of the hsp71 protein M.tuberculosis. Serum from a patient with sputum smear-negative pulmonary tuberculosis, diluted 1:400 was reacted with solid-phase bound peptides using the ELISA method.

15

Figure 7 Shows localization of epitopes within the carboxy-terminal hsp71 sequence. Mean absorbance values from eight adjacent octamer peptides each containing the individual amino acid residues listed on the horizontal scale.

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Sera tested at 1:400 dilution were collected from patients with smear-negative pulmonary tuberculosis (PT, see binding to individual peptides in figure 6), lung disease caused by *M. avium* (AM) and lepromatous leprosy (LL). A-D=localisation of epitope core hexamers.

Figure 8 Shows semi-quantitative evaluation of antibodies to four (A-D) epitopes. Sera diluted 1:400 were collected from patients with smear-positive (full circle) or smear-negative (shaded circle) tuberculosis and from non-tuberculosis controls (open circles) represented by one serum each from lepromatous leprosy, lung mycobacteriosis, Crohn's disease and a healthy subject. All sera were pre-selected to contain high antibody levels to the whole hsp71 antigen. Absorbances (vertical scale) are mean values from adjacent peptides corresponding to each of the four epitopes (bottom horizontal scale).

The sequences of the carboxy terminal regions of *M. tuberculosis* Hsp71 and the corresponding human Hsp70 and *Escherichia coli* dnaK molecules and the Hsp71 of *M. leprae* are set out in Figure 1. Four linear epitopes within the Hsp71 of *M. tuberculosis* are indicated by the

core sequences A to D underlined in Figure 1. It is presently preferred that the peptide of the invention comprises at least one of these linear epitope core sequences or an immunological equivalent thereof.

5 Core sequences A and C, especially the latter, which appear to be more highly specific to M. tuberculosis, are preferred for selective diagnosis of paucibacillary TB in humans. Core sequences B and D appear to be more immunogenic than sequences A and C but have some cross
10 reaction between M. tuberculosis and M. leprae; these core sequences are therefore preferred for more highly sensitive diagnosis of paucibacillary TB where infection by M. leprae can be ruled out on other grounds.

 The linear epitopes may be longer or shorter
15 than the 6-residue core sequences given in Figure 1 but are unlikely to be less than five residues in length.

Accordingly, preferred peptides of the invention comprise at least any five and preferably all six amino acid residues of one of the core sequences indicated in

20 Figure

1. More preferred peptides are octamers containing five or preferably all six of the amino acid residues of one of the core sequences and especially preferred peptides are octamers having the whole core sequence and two other
25 residues corresponding to the two flanking residues at either the N- or C- flanks or one residue at each flank in

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the M. tuberculosis Hsp71 sequence given in Figure 1.

Other preferred peptides of the invention contain a sequence immunologically equivalent to one of the aforementioned linear epitopes, i.e. peptides which cross-react with antibodies detected by peptides containing a linear epitope sequence; more preferably such immunological equivalents contain sequences which cross-react with one of the aforementioned core sequences.

The peptides of the invention may also contain sequences unrelated to Hsp71 or corresponding molecules, at either or both flanks of the core sequence or linear epitope sequence. In a particularly preferred embodiment the peptide comprises a branched polylysine bearing multiple copies of a linear epitope sequence or immunological equivalent thereof.

Peptides of the present invention may be produced by well known peptide synthetic techniques using solid or liquid-phase reactions to assemble the peptide one amino acid residue at a time or by covalently binding preformed sub-sequences, e.g. by linking linear epitope sequences or immunological equivalents thereof to a branched polylysine or other branched structure. Branched polylysines and other suitable branched structures are commercially available or described in the literature and linking of peptides to these may be achieved by conventional techniques.

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Techniques for identifying linear epitopes and immunological equivalents thereof are readily available to those skilled in the art.

5 The peptides of the invention may be used in any conventional immunodiagnostic test technique for detecting antibodies in a body fluid such as blood or serum.

10 The invention further provides a diagnostic test process comprising contacting a peptide as hereinbefore defined with a body fluid sample taken from an individual suspected of having a mycobacterial infection. The peptide may be contacted in the liquid phase or the solid phase, in which case the peptide will be bound to a solid carrier such as the test vessel, for instance a
15 microtitre plate, or latex beads. Binding of antibody from the sample to the peptide may be detected by any conventional technique.

For use in certain diagnostic procedures it may be advantageous to bind the peptide to a detectable label
20 such as an enzyme label, a chromophore, a fluorophore or a radio-isotope or to a solid support. Binding of the peptide to a label or solid supports may be effected by conventional techniques. Peptides bound to such labels and solid supports form a further aspect of the invention.

25 Although it is presently envisaged that the peptides of the invention will be used in in vitro

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diagnostic procedures, it is possible that in vivo techniques will be used. Accordingly, the invention further provides a peptide as hereinbefore defined, optionally bound to a label or solid support, for use in a
5 diagnostic process practised on the human or animal body.

The invention further provides a process for producing a peptide as hereinbefore defined; a process for linking a peptide as hereinbefore defined to a detectable label or solid support; and tests kits containing the
10 peptide optionally bearing a label or linked to a solid support and optionally also comprising buffer, positive or negative control materials, labelling reagents, reagents for detecting a label and other conventional components.

The invention will now be illustrated by the
15 following Examples:-

EXAMPLE 1

INTRODUCTION

In the past, hemagglutinating antibodies to
5 tuberculin PPD were demonstrated in the serum of patients
with active tuberculosis (1,2,3). However, in order to
achieve satisfactory specificity and sensitivity of
serology, antigenic epitopes with both species-specificity
and strong immunogenicity would be required. Recent
10 advances in the molecular definition of mycobacterial
antigens has been exploring possibilities for developing a
clinically useful test for the serodiagnosis of
tuberculosis. In smear-positive (multibacillary)
tuberculosis (TB), this requirement is best met by the 38
15 kDa protein antigen (reviewed in 4). However, in the case
of smear-negative (paucibacillary) disease where laboratory
diagnostic help is most required, the search for an "ideal"
overlap in specificity and immunogenicity within a single
epitope has not as yet been forthcoming.

20 Antibody levels to the 38kDa protein which
represents the most specific and immunogenic antigen in
multibacillary tuberculosis have been found much lower in
smear-negative pulmonary tuberculosis (5,6); nevertheless,
useful diagnostic discrimination of paucibacillary patients
25 with this antigen was achieved using a more sensitive
modification of the competition test (7). Of the other
candidate antigens, about one half of smear-negative
patients had a specific rise of antibodies against the 19kDa
and to a lesser extent to the 14 kDa antigens (6), the

latter being compromised in its discriminatory potential by raised antibody levels in exposed healthy subjects (8). Consequently a combination of tests also involving antibody levels to the M tuberculosis-specific TB78-epitope of the hsp65 protein and lipoarabinomannan have been advocated (9).

In order to expand further the serological evaluation of mycobacterial antigens of defined structure, we examined in this paper antibody levels in patients and controls in respect of the hsp70 antigen. This protein, originally identified by monoclonal antibodies (10) has been sequenced by cloning its gene from the DNA of M. leprae and M tuberculosis (11,12). It belongs to the family of heat shock proteins with broadly conserved structure between prokaryotes and eukaryotes. At this is a strong structural basis for autoimmunization, it was of interest to compare serum antibody levels to both mycobacterial and human hsp 70 molecules in groups of patients and controls.

MATERIALS AND METHODS

20

Serum samples. All 198 tested sera were obtained from patients at the Northwick Park Hospital. (Informed consent was obtained from all patients and controls whose blood was sampled). On the basis of clinical diagnosis, they were classified to groups listed in table 1. Patients with active tuberculosis (group A:90 total) were further classified in respect of pulmonary localization, sputum smear microscopy and bacteriological culture. The clinical parameters of patients with extrapulmonary disease have

previously been described in greater detail (7). Patients with "inactive tuberculosis" (36 total) were represented by patients with radiological evidence of old pulmonary tuberculosis without history of drug treatment (group B "self-healed"), and those with a history of drug treatment (group C "past-treated"). Groups without active or past TB were represented by patients with clinical and radiologically demonstrable lung fibrosis (group D) and by patients with other diseases such as pneumonia, sarcoidosis, bronchiectasis, carcinoma of the lung, viral infections, liver diseases or heart failure (group E). Healthy subjects were age matched adult BCG-vaccinated individuals (group F).

Preparation of antigens: The gene coding the 71kDa protein of M tuberculosis inserted into the PUC8 multicopy plasmid was expressed in TG1 cells and purified by ATP-agarose chromatography (13). The material used in the study (M-hsp70) was prepared and kindly supplied from the WHO IMMTUB/IMMLEP Bank by Dr. Jan van Embden (Bilthoven, The Netherlands).

Recombinant human hsp70 protein was prepared from the stress-inducible hsp70 gene (14) kindly provided by Dr Rick Morimoto, Northwestern University, Evanston, IL., USA. It was subcloned in plasmid PET-3a (15) and expressed in E Coli using bacteriophage T₇ RNA polymerase system for selective high level expression (16). Hsp70 protein was purified from the crude protein lysates (Jindal and Young, in preparation).

The purity of both recombinant hsp70 antigens was

examined by SDS PAGE and Western blot analysis (figure 2). The results showed a single major band of 70 kDa for both the mycobacterial and human recombinant antigens. Only a few minor contaminating bands of lower molecular weight were
5 detected in the H-hsp70 preparation with the human serum which reacted with several proteins in the crude E coli extract.

Table 1. Clinical Classification Of Patients

Group*	Clinical Diagnosis	Bacterial Status		
		S ⁺ C ⁺	S ⁻ C ⁺	S ⁻ C ⁻
A	Pulmonary	30 ⁰	14	9
	Active TB Extrapulmonary	8	9	20
B	Inactive TB: self-healed	0	0	21
C	past-treated	0	0	15
D	Lung fibrosis	0	0	5
E	Other diseases	0	0	31
F	Healthy subjects	0	0	20

* - Corresponding with designations in figure 12

⁰ - Number of patients

S = smear microscopy (+ or -)

C = culture of *M.tuberculosis* (+ or -)

Enzyme Linked Immunoassay (ELISA). Polyvinyl microtitre plates (Nunc-Immuno plate MaxiSorp F96) were incubated with 50 μ l of either M-hsp70 or H-hsp70 at concentration of 1 μ g/ml or 2 μ g/ml respectively at 4°C overnight. After

5 washing once with phosphate-buffered saline containing 0.05% (w/v) "Tween 20" (PBST), the wells were incubated with 200 μ l of 2% dried milk (w/v) in PBST (PBSTM) at 37°C for 1 hr. Four fourfold dilutions of human sera (1/50-1/3200) using 5 μ l volume were added to duplicate wells and incubated at

10 37°C for 2 h. After 5 washings with PBST, 50 μ l of affinity purified goat antihuman IgG peroxidase conjugate (diluted 1/1000) was added to each well and the plates were further incubated for 1 h. After washing with PBST, 75 μ l tetramethyl benzidine plus hydrogen peroxide in citrate

15 buffer at pH 5.1 was added to each well; the reaction was stopped in 10 min. time with 25 μ l per well 0.5 mol/L sulphuric acid. The absorbance at 450 nm was read in the "Titertek Multiskan II" spectrophotometer. Antibody titers were expressed as the dilution of serum giving 30% of the

20 plateau binding value of a standard positive serum (ABT₃₀).

Statistical analysis: Means and standard deviations of the log₁₀ values of antibody titers were calculated for each group of patients and controls. Groups were compared with

25 each other by the student t-test for unpaired values. The proportion of patients with increased antibody levels in groups A-D was calculated with 98% specificity of the cut-off limit in comparison with values in group E.

RESULTS

Patients with diseases other than TB (group E) had antibody levels to both M-hsp70 and H-hsp70 significantly ($P < .001$) elevated above those found in healthy individuals (group F) (Figure 3). Therefore antibody levels in group E rather than those of the healthy controls (group F) were used as the "baseline" for evaluating the extent and significance of rise in antibody levels in groups with either active or inactive TB.

Patients with active tuberculosis (group A, Figure 3) had IgG antibody titers to M-hsp70 (mean $ABT_{30} 3.2 \pm 0.4$) significantly higher ($P < .001$) than that of non-TB patients (group E) (mean $ABT_{30} 2.3 \pm 0.4$). When taking the anti-M-hsp70 \log_{10} titer 2.75 as the "cut-off" value 78% out of the 70 tested patients with active TB and significantly raised antibody levels. Corresponding evaluation of anti-H-hsp70 antibody levels using the \log_{10} titer 2.5 as the discriminatory cut-off value in respect of group E, produced a somewhat lower, 50% positivity rate in patients with active TB.

Analysis of titers in groups of patients with "inactive tuberculosis" in comparison with group E demonstrated that antibody levels have been increased to a significantly greater extent in self-healed (group B: 80% for M-hsp70 and 67% for H-hsp70) than in the past treated (group C: 53% for M-hsp70 and 47% for M-hsp70) individuals. In view of the large individual variations, we examined the possible association of antibody levels with several clinical

criteria such as age, race, duration of illness, chest X-ray, weight loss, fever and history of tuberculosis. However, none of these parameters were significantly correlated with antibody levels.

5 Interestingly, the clinical evaluation revealed that both anti-M and anti-H hsp70 titers were the highest in five patients of group B (boxed in figure 3) who were on basis of prior independent clinical and radiographic evaluation considered to be at risk of relapse and were
10 therefore put on "secondary" rifampicin/isoniazid chemoprophylaxis. However, irrespective of the tuberculosis process, uniformly high antibody levels were also observed in all five tested patients with lung fibrosis (group D). Since most of the non-tuberculosis pulmonary and
15 extrapulmonary diseases had "chronic inflammation" but did not have raised antibody levels, it would seem that anti-hsp70 antibodies are not stimulated merely by inflammation but rather represent a marker of the next stage, ensuing fibrosis.

20 It was of further interest to break down the antibody levels within the group with active TB (column A in figure 3) in relation to bacterial smear and culture positivity. The values expressed in Figure 4 showed no significant differences between the multibacillary (S⁺C⁺),
25 intermediary (S⁺C⁺) and paucibacillary (S⁺C⁻) sub-groups in respect of either anti-M or anti-H- hsp70 antibody titers.

Comparison of antibody titers to M- and H- hsp70 antigen in individual patients showed highly significant correlation coefficients for all tested groups (mean R value

= 0.66). Nevertheless, it was noted that patients with active TB, had a relatively high ratio of titres to mycobacterial compared to human hsp70 (M/H ratio 4.2). The form of tuberculosis of the 11 patients with > 10 M/H ratio was pulmonary (3x), lymph node (4x), bone & skin (3x) and pericardial (1x). In contrast, a higher proportion of anti-human hsp70 titres (M/H ratio = 1.8) was found in patients with lung fibrosis and other non-TB diseases (figure 5).

10 DISCUSSION

Advances in the structural definition of mycobacterial antigens enabled a systematic study of specificity of the immune repertoire (4). Since, the relationships between the structure of antigenic molecules and their immune functions following infection with pathogenic mycobacteria are being established. The antibody repertoire of mice has been found to be distinctly more selective during infection with M. tuberculosis than following artificial immunization procedures (17). Hence, potential serodiagnostic antigens with high immunogenicity and specifically for the infecting pathogen need to be selected on the basis of detailed knowledge of the human immune system to pathogenic mycobacterial infection. The hsp70 protein, chosen for the present study, represents a well known major stress protein which is highly inducible under stress conditions (e.g. heat shock) and is one of the HLA linked hsp70 genes (18). Proteins of this family "chaperone" the folding and transport of intracellular

polypeptides. The 71kDa protein of M tuberculosis has biochemical and functional features consistent with other members of the hsp family (13) and has been found in supernatants from short-term culture of tubercle bacilli
5 (19). The extracellular location may enable contact of the molecule in its native conformation with B cells which is a requirement for effective stimulation of antibody formation.

Despite the greatly conserved amino acid sequence composition of hsp70, it is of interest that existing
10 monoclonal antibodies have been found restricted in binding mainly to pathogenic mycobacteria of the M. tuberculosis complex. M. leprae and M. scrofulaceum (10). Following infection with Schistosoma mansoni, antibodies also exhibited only limited cross-reactivity in respect of
15 species specific epitopes of hsp70 and arose early in the course of infection (20,21). However, sera from humans infected with malaria contained autoantibodies which reacted with human hsp70 (22). Increased antibody levels to mycobacterial hsp70 have been found in sera of patients with
20 active pulmonary tuberculosis (23, 24), but these studies did not show individual values and did not evaluate antibody levels in relation to the stage of disease.

The results in this study demonstrated significant elevation of anti-hsp 70 antibody levels in the majority
25 (78%) of patients with active pulmonary TB. Furthermore, about 40% (based on the ratio between titres) of the antibodies appeared to be cross-reactive with the human hsp70 molecule. Despite its highly conserved aminoacid structure, the stimulation of anti-hsp70 antibodies appears

to be strongly associated with the tuberculosis infection since patients with a broad range of other diseases (excluding tuberculosis) had much less raised antibodies that TB patients, when compared with healthy controls. In this respect the anti-hsp70 response follows a quite different pattern than that to hsp65 which showed a very broad range of titres in non-TB controls which prevented any meaningful discrimination from tuberculosis patients (6). However, the persistence of raised hsp70 antibody levels for several years following active tuberculosis, which is known to apply also to other antigenic specificities, represents a caveat for diagnostic consideration.

The most striking finding has been the lack of any difference in the serological response to hsp70 between multibacillary and paucibacillary cases of tuberculosis. This finding is in contrast with the much more pronounced response of smear-positive TB patients to the previously examined M. tuberculosis specific epitopes (5,6). Obviously there may be a limitation for diagnostic application due to the reported elevation of anti-hsp70 levels in some autoimmune diseases such as systemic lupus erythematosus and rheumatoid arthritis (24,25). However, in autoimmune patients and also in the inactive TB cases, the antibodies appear to be predominantly cross-reactive with human hsp70. On the other hand it is encouraging that the anti-mycobacterial titres were approximately four times higher in the diagnostically difficult smear-negative and extrapulmonary cases as often as in smear-positive pulmonary TB. In fact, of the 11 sera with < 10 M/H ratio, 73% were

from patients with extrapulmonary TB. Hence, there would
would be scope for improving further the mycobacterial
specificity of the serology by using synthetic peptides or
truncated recombinant segments (26) from the carboxy-
5 terminal portion of the molecule which contains amino acid
sequences with the most pronounced specificity for
M.tuberculosis (12).

The mechanisms, which cause the apparently unique
immunogenicity of hsp70 in the early stages of tuberculosis
10 need further study. Proliferative responses of peripheral
blood CD4 T cells to mycobacterial hsp70 were observed to be
somewhat stronger than those to hsp65 (27) and CD8 T cell
clones of hsp70 specificity were obtained from the
peripheral blood of tuberculosis patients (28). The
15 reported role of host hsp70 in processing of antigenic
peptides (29) and expression on the surface of stimulated
macrophages (30) could play a role for enhancing antibody
responses to low levels of antigenic stimulation. The
reported induction of synthesis of bacterial hsp70 together
20 with other stress proteins during intracellular replication
(31) could promote bacterial persistence via the ATP binding
activity or by the special genetic regulatory elements (12)
of hsp70. The pronounced serological response to hsp70 may
indeed be signalling a distinct yet unrecognised function of
25 this protein in the pathogenic host-parasite relationship.

REFERENCES FOR EXAMPLE 1

1. Middlebrook G, Dubos R, Specific serum
agglutination of erythrocytes sensitised with
5 extracts of tubercle bacilli. J Exp Med
1948;88;521-528.
2. Boyden SV. The adsorption of proteins on
erythrocytes treated with tannic acid and
10 subsequent hemagglutination by antiprotein sera. J
Exp Med 1951;93;107-120.
3. Turcotte R, Freedman SO, Sehon AH. A new
hemagglutination procedure for the diagnosis of
15 active tuberculosis. Am Rev Resp Dis 1963;88;725-
728.
4. Ivanyi J, Sharp K, Jackett P, Bothamley G.
Immunological study of the defined constituents of
20 mycobacteria Springer Semin Immunopathol
1988;10:279-300.
5. Bothamley G, Udani P, Rudd r, Festenstein F, Ivanyi
J. Humoral response to defined epitopes of tubercle
25 bacilli in adult pulmonary and child tuberculosis.
Europ J Clin Microbiol Infect Dis 1988;7:639-545.
6. Jackett PS, Bothamley GH, Batra HV, Mistry A, Young
DB, Ivanyi J. Specificity of antibodies to

immunodominant mycobacterial antigens in pulmonary tuberculosis J Clin Microbiol 1988;26:2313-2318.

7. Wilkins EGL, Ivanyi J. The potential value of serology for the diagnosis of extrapulmonary tuberculosis. The Lancet 1990;336:641-644.
8. Hoepfner VH, Jackett PS, Beck JS, Karjirto T, Grange, Ivanyi J. Appraisal of the monoclonal antibody based competition test for the serology of tuberculosis in Indonesia, Serodiagnosis and Immunotherapy 1987;1:69-77.
9. Bothamely G.H, Rudd R, Festenstein F, Ivanyi J. Clinical value of the measurement of M tuberculosis-specific antibody in pulmonary tuberculosis. Chest 1991; in press.
10. Britton WJ, Hellqvist L, Basten A., Raison R.L. Mycobacterium leprae antigens involved in human immune responses. Identification of four antigens by monoclonal antibodies. J Immunol 1985;135:4171-4177.
11. Garsia RJ, Hellqvist L, Booth RJ, Radford AJ, Britton WJ, Asbur, L, Trent RJ Basten A. Homology of the 70-kilodalton antigens from Mycobacterium leprae and Mycobacterium tuberculosis 71-kilodalton antigen and with the conserved heat shock protein

70 of eucaryotes. Infect Immun 1989;57:204-212.

12. Lathihgra R, Alexander W. Stover K, Coadwell J,
Young R, Young D. Genetic Characterization of the
5 dnaK-grpE-dnaJ heat shock gene cluster in
Mycobacterium tuberculosis in press.
13. Mehlert A, Young D.B. Biochemical and antigenic
characterisation of the Mycobacterium tuberculosis
10 71 kD antigen, a member of the 70kD heat-shock
protein family. Mol Microbiol 1989; 3:125-130.
14. Hunt C, Morimoto RI. Conserved features of
eukarotic HSP70 genes revealed by comparison with
15 the nucleotide sequence of human hsp70. Proc.
Natl. Acd. Sci. USA 1985; 82:6455-6459.
15. Rosenberg AH, Lade BN, Cui DS, Lin S-W, Dunn J,J
Studier FW. Gene 1987;56:125-135.
- 20 16. Studier, FW, Moffatt BA. Use of bacteriophage T₇ RNA
Polymerase to direct selective high level
expression of cloned genes. J Mol Biol
1986;189:113-130.
- 25 17. Brett SJ, Ivanyi J. Genetic influences on the
immune repertoire following tuberculosis infection
in mice. Immunology 1990; 71:113-119.

18. Sargent CA, Dunham I, Trowsdale J, Cambell RD,
Human major histocompatibility complex contains
genes for the major heat shock protein HSP70. Proc
Natl Acad Sci USA 1989;86;1968-1972.
- 5
19. Abou-Zeid C, Smith I, Grange JM, Ratliff TL Steele
J, Rook GA.W. The secreted antigens of M.
tuberculosis and their relationship to those
recognised by the available antibodies. J. Gen
10 Microbiol 1988;134;531-538.
20. Hedstrom R, Culpepper J, Harrison RA, Agabian N,
Newport G, A major immunogen in Schistosoma Mansoni
infections is homologous to the heat-shock protein
15 Hsp70. J Exp Med 1987;165:1430-1435.
21. Newport GR, Hedstrom RC, Kallestad J, Tarr P,
Klebanoff S, Agabian N. Identification, molecular
cloning and expression of a schistosome antigen
20 displaying diagnostic potential. Am. J. Trop Med
Hyg 1988; 540:540-546.
22. Mattei D, Scherf A, Bensaude O, da Silva LP, A
heat shock-like protein from the human malaria
25 parasite Plasmodium falciparum induces
autoantibodies. Eur J Immunol 1989;19:1823-1828.
- 23 Britton WJ, Hellqvist L, Basten A, Inglis AS,
Immunoreactivity of a 70 kDa protein purified from

Mycobacterium bovis bacillus Calmette-Guerin by monoclonal antibody affinity chromatograph J Exp Med 1986;164:695-708.

- 5 24. Tsoulf, G, Rook GAW, Bahr, GM, Sattar, MA,
Behbehani K, Young DB, Mehlert A, van-Embden JDA,
Hay FC, Isenberg DA, Lydyard PM. Elevated IgG
antibody levels to the mycobacterial 65-kDa heat
shock protein are characteristic of patients with
10 rheumatoid arthritis. Scand J Immunol 1989;30:519-
527.
25. Minota S, Cameron B, Welch WJ, Winfield JB,
Autoantibodies to the constitutive 73-kD member of
15 the hsp70 family of heat shock proteins in systemic
lupus erythematosus. J Exp Med 1988;168:1475-1880.
26. Mehra V, Sweetser D, Young RA, Efficient mapping of
protein antigenic determinants. Proc Natl Acad Sci
20 USA 1986;83:7013-7017.
27. Adams E, Garsia RJ, Hellqvist L, Holt P, Basten A T
cell reactivity to the purified mycobacterial
antigens p65 and p70 in leprosy patients and their
25 household contacts. Clin Exp Immunol 1990;80:206-
212.
28. Rees A, Scoging A, Mehlert A, Young DB, Ivanyi J.
Specificity of proliferative response of human CD8

clones to mycobacterial antigens. Eur J Immunol
1988;18:1881-1887.

29. Vanbuskirk A, Crump BL, Margoliash E, Pierce SJ. A
5 peptide protein having a role in antigen
presentation is a member of the Hsp70 heat shock
family. J Exp Med 1989: 170:1799-1809.
30. Jarjour W, Tsai V. Woods V, Welch W, Pierce S, Shaw
10 M, Mehita H, Dillmann W, Zvaifler N, Winfield J.
Cell surface expression of heatshock proteins.
Arthritis Rheum 1989:32-S44.
31. Buchemeier N.A, Heffron F, Induction of Salmonella
15 stress proteins upon infection of macrophages.
Science 1990;248:730-732.

EXAMPLE 2

20 INTRODUCTION

The specificity of the immune repertoire in
mycobacterial infections has been of interest from several
aspects of the immunopathogenesis as well as for potentials
25 towards diagnostic and vaccine development. Serological
studies suggested that the antibody response in respect of
species-specific protein antigens is restricted to
relatively few constituents represented by the 38kDa, 19kDa
and 14kDa antigens in tuberculosis and by the 35 kDa antigen

in leprosy (1). Antibody levels to all these antigens have been found most pronounced in patients with multibacillary forms of the disease which are clinically the most apparent and can be confirmed by bacteriological techniques.

5 However, the sensitivity of detection of paucibacillary infection i.e. smear/culture negative tuberculosis using single or several antigenic probes has been less satisfactory (2-4).

A new important advancement in this direction has
10 been achieved recently, by the demonstration that antibody levels to the mycobacterials hsp71 protein have been elevated to the same extent in patients with either smear-positive or smear-negative tuberculosis (5). The hsp71 antigen, originally identified by monoclonal antibodies is a
15 stress protein constituent of mycobacterial (6-8). In view of this new opportunity for early serological diagnosis of paucibacillary disease it has been of further interest to identify those hsp71 epitopes which are both immunodominant as well as M.tuberculosis-specific. The need for focusing
20 on the selection of species specific epitopes has been mandatory, to avoid the detection of cross-reactive antibodies to those parts of the hsp70 molecule which carries extensive structural homology with the corresponding proteins contained in all microbial and eukaryotic cells
25 (7). Thus, we had chosen for epitope scanning the carboxy-terminal part of the molecule which has been found most polymorphic, in alignment of the hsp70 sequences from mycobacteria and other species(9).

MATERIALS AND METHODS

Peptides

The general net synthesis of the peptides purchased from
5 Cambridge Research Biochemicals (CRB) (Cambridge, England)
was comprised of 94 peptides based on 101 residues on the
basis of the sequence of hsp71 of *M. tuberculosis* residues
509-621 (10). Non-cleavable octapeptides, overlapping by
one residue starting at residue 509 (VRNQAETL) and ending at
10 residue 621 (GRCPRLG) were synthesised in solid-phase using
the Geysen technique (11) on specially moulded, high-density
polyethylene pins which are assembled into a plastic holder
designed to hold 96 pins in the format and spacing of a
microtitre plate. The tips of the pins have been chemically
15 derivatised to provide functional groups to which amino
acids may be coupled in an appropriate form in peptide
synthesis. All peptides have received N-terminal
acetylation. Positive and negative control peptide
sequences were synthesised on the same block as the test
20 peptides to give an indication of the sensitivity being
achieved in the ELISA assay.

Sera

Selected sera which had previously been tested and all found
as a high titre (1/2000-1/4000) to recombinant mycobacterial
25 hsp70 (5) were collected from 2 cases of smear-positive
pulmonary tuberculosis, 6 cases of smear-negative
tuberculosis, and 1 case each of lepromatous leprosy, non-
tuberculosis pulmonary disease of proven infection with *M.*
avium, Crohn's disease and one healthy subject.

ELISA

This was performed following the recommended procedure from CRB. Reactions at the tips of the pins are carried out in polypropylene plates with wells in a standard 8 by 12 matrix matching the pin spacing. Briefly, the pins were precoated by dispensing 200 μ l of pre-coat buffer (2% w/v bovine serum albumin), 0.1% v/v Tween 20, 0.01 phosphate buffered saline pH 7.2) into each well of a microtitre plate. The pins were placed in the wells and incubated for 60 min at 20°C on a shaker at 100 rpm. Without washing, the pins were incubated overnight at 4°C in a microtitre plate containing 175 μ l/well of human serum diluted 1/400 in pre-coat buffer. The pins were removed from the primary antibody preparation and washed four times 10 min each in a bath of 0.01 M PBS (pH 7.2) at room temperature with agitation. Then, the pins were placed in a microtitre plate containing 175 μ l/well of peroxidase labelled goat anti-human IgG diluted 1/1000 in conjugated diluent (1% v/v sheep serum, 0.1% v/v Tween 20, 0.1% w/v sodium caseinate and 0.01 M PBS pH 7.2) and incubated at 20°C for 60 min with agitation. The pins were washed as above and placed in a microtitre plate containing 150 μ l/well of substrate solution, 0.5 mg/ml of 2,2'-azino-di[3-ethyl-benthiazoline sulfonate] in citrate buffer pH 4.0 and 0.01% w/v hydrogen peroxide. The reaction was stopped by removing the pins from the plate after 30 min and immediately the optical density was measured at 405 nm in a Titertek "Multiscan MCC II" spectrophotometer.

RESULTS

Altogether 13 sera were successively tested for binding to 94 overlapping peptides presented on a single 96-pin polyethylene block. There was no indication of any deterioration in the binding activities of peptides as judged by closely similar binding values to two pins containing control peptides. Optical density values from the ELISA assay of one representative serum from a patient with smear-negative pulmonary tuberculosis showed elevated antibody binding to several discretely separated clusters of adjacent peptides (figure 6). Four areas of peptides, based on sequences towards the carboxy-terminus and with the most consistent localisation of increased binding had been designated as A (peptides 33-41), B (peptides 44-52), C (peptides 56-67) and D (peptides 74-85). Elevated binding values were also observed to peptides located further towards the amino-terminus; however, these presumably cross-reactive epitopes were omitted from evaluation. Peptide No. 77 showed low binding with all tested sera which we considered to be due to defective peptide structure or presentation in view of the consistent performance of adjacent peptides on both sides from peptide 77.

In order to evaluate the contribution of individual amino acids to the epitope structure, we have calculated the mean absorbance values for each residue from all 8 overlapping octapeptides containing the respective amino acid. This analysis was applicable to sequences, starting from the leucine at position 516, which represented the last

residue in peptide number 1. The results shown in Figure 7 demonstrate the values obtained from testing one serum each from patients with pulmonary tuberculosis (PT), calculated from values shown in figure 1), lung disease caused by M. avium (AM) and with lepromatous leprosy (LL). Antibody binding activities towards individual residues closely overlapped in their localisation and were clearly pronounced for epitopes B and D with each of the three tested sera. In contrast, binding to epitopes A and C identified with serum PT was not significant with the AM and LL sera. This individual variation in the binding patterns between sera supports the view that the four identified epitopes are of distinct specificity.

The core structures of each of the four identified epitopes have been allocated to six residues graded with the highest mean OD values. They are represented by sequences DAAVAE for epitope A, WRIGYF for epitope B, GEAGPG for epitope C and CVTGHW for epitope D (Figure 1). Alignment of these hexamers within the sequence of the hsp71 carboxy terminus from M. tuberculosis (10), with corresponding ssequences of M.leprae hsp70 (9), E.coli dnaK, (12) and of the human HSP70-1 gene (13) revealed, that the core of epitope A is overlapping in 5 out of the 6 core residues with the M.leprae sequence. In contrast, epitopes D, B and C shared only 2, 1 or no residue respectively with M.leprae. In view of the shared sequence of epitope A, it was surprising to find a lack of binding by the LL serum, selected with a high titre to several mycobacterial antigens. None of the four epitopes had significant

homology with either dnaK or human HSP70-1 structure.

Predictive epitope evaluation of the carboxy-terminal sequence by the "Protean" programme (Proteus Biotechnology Ltd, Manchester, UK; kindly performed by J. Edwards at
5 Peptide Technology Ltd. Sydney, Australia) allocated the highest score only to a single broad stretch of residues 565-591 which contains the epitope C (571-576) sequence.

A semi-quantitative evaluation of antibodies in the 13 tested sera against individual epitopes has been done by
10 calculation of mean OD values from the 7 - 11 peptides, corresponding to each of the four epitopes. The results represented in Figure 8, demonstrate significantly higher antibody binding to each of the four epitopes in sera from tuberculosis patients when compared with non-tuberculosis
15 controls. This difference was more significant for epitopes C and D ($p < 0.001$) than for epitopes A and B ($p < 0.01$). However, it should be remembered, that the serum from the healthy subject and the other 3 controls were purposely selected to contain high antibody binding to fulfil the
20 topographic aims of the study. Hence, much better discrimination would be expected between TB patients and randomly selected non-tuberculosis patients and healthy controls.

DISCUSSION

25

The immunogenicity of protein antigens in terms of antibody formation is largely based on the recognition of epitopes displayed on native molecules by the immunoglobulin receptors of B cells (14). Indeed, the epitopes which react

with the bulk of antibodies in patients with tuberculosis are of conformational nature (1, 15, 16). Furthermore, the need for direct contact with B cells by the native antigens is corroborated by the fact that both 38kDa and 19kDa
5 antigens are lipoproteins, secreted from live mycobacteria (17) and resulting in the highest antibody titres in multibacillary disease (2,3). In the paucibacillary stage of infection however, the presentation of those antigens of
10 intracellularly replicating mycobacteria which had been processed within macrophages would stimulate an antibody repertoire of different specificity, directed predominantly to linear epitopes. This view is supported by the recent demonstration that antibody levels to the mycobacterial hsp71 antigen have been raised equally in sera from both
15 bacteriologically positive and negative cases of tuberculosis (5). Similarly, anti-hsp70 antibodies were detected in the early stages of schistosomiasis (18). The pronounced immunogenicity of hsp71 during paucibacillary infection could be attributed to enhanced stress-induced
20 synthesis during intracellular replication (19), or to surface expression (20,21) enabling recognition by the Ig receptors of B cells or merely to earlier priming by multiple cross-reacting commensal microbial organisms and autologous hsp70. Although the hsp71 protein is a
25 predominantly intracellular constituent (8), its presence in media from short-term mycobacterial cultures (22) indicates some form of secretion from live bacteria which could be contributory to its immunogenicity for B cells.

In view of the extensive homology in the primary

structure of prokaryotic and eukaryotic hsp70 proteins (9), antibodies are likely to be produced against many cross-reactive epitopes of the hsp71 antigen. Indeed, elevated serum antibody levels to both microbial and human hsp71 have
5 been found in a variety of infectious, chronic inflammatory and autoimmune diseases (5, 6, 23-26). However, it is notable, that antibodies produced following infection with *Schistosoma mansoni* were found to be predominantly directed against species-specific epitopes of hsp70 (29). The
10 present peptide-scan analysis of the most polymorphic carboxy-terminal part of the hsp71 sequence identified four distinct linear epitopes. Localisation of these four epitopes was highly consistent with all 13 tested human sera without demonstrable microheterogeneity in the epitope
15 structure which could have been revealed by antibodies selected from the broad human B cell repertoire.

Despite the limitation in the number of cycles in which the peptide-pins can be re-used, the analysis of a selected collection of sera allowed the unequivocal
20 localiation of four core epitope sequences. Future bulk synthesis of these peptides will enable more extensive serological evaluation of the identified epitopes. It is possible, that the *M.tuberculosis* epitope aligned sequences from other microbial and eukaryotic species may also carry
25 immunogenic epitopes. Their potential as species-specific probes for the involvement of hsp70 could be valuable in the study of immunopathogenesis of various infectious diseases or autoimmunity and possibly for immunodiagnostic applications.

REFERENCES for Example 2

1. Ivanyii J., Sharp K., Jackett P., and Bothamley G.
(1988) Immunological study of the defined
5 constituents of mycobacteria Springer Seminars in
Immunopathology 10:279-300.
2. Bothamley G., Udanl P., Rudd R., Festenstein F.,
Ivanyi J. (1988) Humoral response to defined
10 epitopes of tubercle bacilli in adult pulmonary and
child tuberculosis Europ J Clin Microbiol 7: (5)
639-545.
3. Jackett P.S., Bothamley G.H., Batra H.V., Mistry
15 A., Young D.B and Ivanyi J. (1988) Specificity of
antibodies to immunodominant mycobacterial antigens
in pulmonary tuberculosis. J. Clin Microbiol 26:
2313-2318.
- 20 4. Wilkins E.G.L., and Ivanyi J. (1990) the potential
of serology for the diagnosis of extrapulmonary
tuberculosis. The Lancet 336:641-644.
5. Elsaghler A.A.F., Wilkins E.G.L., Mehrotra P.K.,
25 Jindal S. and Ivanyi J. (1991) Elevated antibody
levels to stress protein hsp70 in smear-negative
tuberculosis. Immunol & Inf. Diseases. In press
(Example 1).

6. Britton, W.J., Hellqvist, L., Basten, A and Inglis A.S., (1986) Immunoreactivity of a 70 kDa protein purified from *Mycobacterium bovis* Bacillus Calmette-Guerin by monoclonal antibody affinity chromatograph J. Exp. Med 164: 695-708.
- 5
7. Garsia R.J., Hellqvist L., Booth R.J., Radford A.J., Britton W.J., Astbury L., Trent R.J., and Basten A. (1989) Homology of the 70-kilodalton Antigens form *Mycobacterium leprae* and *Mycobacterium tuberculosis* 71-kilodalton Antigen and with the Conserved Heat Shock Protein 70 of Eucaryotes. Infection and Immunity, 57:204-212.
- 10
8. Mehlert A., and Young D.B. (1989) Biochemical and antigenic characterisation of the *Mycobacterium tuberculosis* 71 kD antigen a member of the 70kD heat-shock protein family. Molecular Microbiology. 3:(2):125-130.
- 15
9. McKenzie K.R., Adams E., Britton W.J., Garsia R.J. and Basten A. (1991) Sequence and immunogenicity of the 70-kDa heat shock protein of *Mycobacterium leprae*. J. Immunol 147:312-319.
- 20
10. Lathihgra R., Alexander W., Stover K., Coadwell J., Yound R., and Young D. (19-- Genetic characterisation of the dnaK-grpE heat shock gene cluster in *Mycobacterium tuberculosis*. In press.
- 25

11. Geysen HM., Meloen RH., and Bartelling SJ. (1984)
Use of peptide synthesis to probe viral antigens for
epitopes to a resolution of a single amino acid.
Proc. Natl. Acad. Sci. USA 81: 3998-4002.
- 5
12. Bardwell J. and Craig E. (1984) Major heat shock
genes for *Drosophila* and the *Escherichia Coli* heat-
inducible *dnaK* gene are homologous. Proc. Natl
Acad. Sci. USA. 81:848-852.
- 10
13. Jindal S., Dudani A.K., Singh B., et al (1989)
Primary structure of a human mitochondrial protein
homologous to the bacterial and plant chaperonins
and to the 65-kilodalton mycobacterial antigen.
Mol Cell Biol 248:730-732.
- 15
14. Lanzavecchia A. (1990) Receptor-Mediated antigen
uptake and its effect on antigen presentation to
class II-restricted T lymphocytes. Annu. Rev.
Immunol. 8: 773-793.
- 20
15. Andersen A.B. and Hansen E.B. (1989) Structure and
mapping of antigenic domains of protein antigen b,
a 38,000-molecular-weight protein of *Mycobacterium*
tuberculosis. Infect Immun. 57: 2481-2488.
- 25
16. Ashbridge K.R., Prestidge R.K., Booth R.J. and
Watson J.D. (1990) The mapping of a serodominant
region of the *Mycobacterium tuberculosis* 19

kilodalton antigen. J. Immuno. 144:3137-3142.

17. Young D.B. and Garbe T.R. (1991) Lipoprotein
antigens of Mycobacterium tuberculosis Res.
5 Microbiol. 142:55-65.
18. Hedstrom R., Culpepper J., Harrison R.A., Agabian
N., and Newport G., (1987) A major immunogen in
Schistosoma Mansoni infections is homologous to the
10 heat-shock protein Hsp-70. J. of Exp. Med. 165:
1430-1435.
19. Buchemeier N.A., and Heffron F., (1990) Induction
of Salmonella stress proteins upon infection of
15 macrophages. Science 248:730-732.
20. Vanbuskirk A., Crump B.L. Margollash E., and Pierce
S. J. (1989) A peptide binding protein having a
role in antigen presentation is a member of the
20 Hsp70 heat shock family. Exp Med. 170:1799-1809.
21. Jarjour W., Tsal V., Woods., Welch W., Pierce S.,
Shaw M., Mehita H., Dillmann W., Zvalfler N.,
Winfield J. (1989) Cell surface expression of
25 heatshock proteins. Arthritis Rheum 32-S44.
22. Abou-Zeid C., Smith I., Grange J.M., Ratliff T.L.,
Steele J., and Rook G.A.W. (1988) The secreted
antigens of M. tuberculosis and their relationship

to those recognised by the available antibodies.
J. Gen Micro. 134:531-538.

23. Minota S., Cameron B., Welch W.J. and Winfield J.B.
5 (1988) Autoantibodies to the constitutive 73-kD
member of the hsp70 family of heat shock proteins
in systemic lupus erythematosus J. Exp. Med. 168:
1475-1880.
- 10 24. Tsoulfa G., Rook G.A.W., Bahr G.M., Sattar M.A.,
Behbehani K., Yound D.B., Mehlert A., van-Embden
J.D.A., Hay F.C., Isenberg D.A., and Lydyard, P. M.
15 (1989) Elevated IgG antibody levels to the
mycobacterial 65-kDa heat shock protein are
characteristic of patients with rheumatoid
arthritis. Scand. J. Immunol. 30: 519-527.
25. Mattei D., Scherf A., Bensaude O., and deed of
assignment Silva L.P. (1989) A heat shock-like
20 protein from the human malaria parasite Plasmodium
falciparum induces autoantibodies Eur. J. Immunol.
19 1823-1828.
26. Elsaghier A., Prantera C., Bothamely G., Wilkins
25 E., Jindal S. and Ivanyi J. (1991) Disease
association of antibodies to human and
mycobacterial hsp70 and hsp60 stress proteins.
Submitted for publication.

27. Newport G.R., Hedstrom R.C., Kallestad J. Tarr P.,
Klebanoff S., Agablan N. (1988) Identification,
molecular cloning and expression of a schistosome
antigen displaying diagnostic potential. Am J.
Trop. Med. Hyg 540:540-546.
- 5.

Claims

1. A peptide comprising a sequence the same as or immunologically equivalent to a linear epitope of the carboxy terminal region of Micobacteria tuberculosis Hsp71 or a
5 corresponding heat shock protein or any other mycobacterial pathogen.

2. A peptide according to claim 1 comprising a sequence the same as or immunologically equivalent to a linear epitope of the carboxy terminal of M.tuberculosis
10 Hsp71.

3. A peptide according to claim 1 containing a sequence comprising at least five residues of one of the following linear epitope core sequences:

15 DAAVAE
WRIGYF
GEAGPG
CVTGHW

4. A peptide according to claim 3 containing a sequence:

20 DAAVAE
WRIGYF
GEAGPG
CVTGHW

5. A peptide according to claim 4 containing one of the following sequences:

5 KVDAAVAE
VDAAVAEA
DAAVAEAE
GTWRIGYF
TWRIGYFG
WRIGYFGH
GDGEAGPG
10 DGEAGPGV
GEAGPGVA
SGCVTGHW
GCVTGHWR
CVTGHWRC

15 6. A peptide according to claim 1 or claim 2 containing a sequence immunologically equivalent to a sequence according to any one of claims 3 to 5.

7. A peptide according to any preceding claim bound to a solid carrier or a detectable label.

20 8. A peptide according to claim 7 bound to an enzyme label, cromophore, fluorophore or radio-isotope.

9. A peptide according to any preceding claim for use in a diagnostic process practised on the human or animal body.

10. A diagnostic test process comprising contacting a peptide according to any one of the preceding claims with a body fluid sample taken from an individual suspected of having a microbacterial infection.

11. A diagnostic test kit comprising a peptide according to any one of claims 1 to 9 optionally bearing a label or linked to a solid support and optionally also comprising buffer, positive or negative control materials, labelling reagents, reagents for detecting a label and other conventional components.

Fig.1.

Hu: A K . A L . S Y A F N M . S A . . D E G L K
 Ec: T . . . G D H . L H S . R . Q . E . A G D K
 Lp: T
Tb: V R N Q A E I L V Y Q I E K F V K E Q R E A 530

- . - . . I S . A D K K . . L D K C Q . V I S W
 L P A D D K T A I E S A L T A L E T . L K G E -
 . N . . R E T A
E G G S K V P E D I L N K V D A A V A E A E G - 554

=====A=====

L D A N T L - A E K D E F E H K R K E L E L Q V
 - D K A A . E A K M Q E L A Q V S Q K L M E I .
 L . G T D . - S A I K S A M E K L G Q D S Q A L
 - G I W R I - G Y F G H Q V G D G E A G P G V A 578

=====B=====

=====C=====

C N P I I S G L Y Q G A G G - - P G P G G F G A
 Q Q Q H A Q Q Q T A G A D A S A N N A K D - - -
 . Q A I Y E A T Q A A S K . G . E A S A . G G S
G S G A S D L R S S S G C V I G H W R C P - - - 602

=====D=====

Q G P K G - G S G S G P T I E E V D -
 - - - D D V V D A E F E E V K D - K K
 N S T D D V L T . . W S T T N G S P K
 - - - - - P R R R A G R C P P R L G 621

Fig. 2.

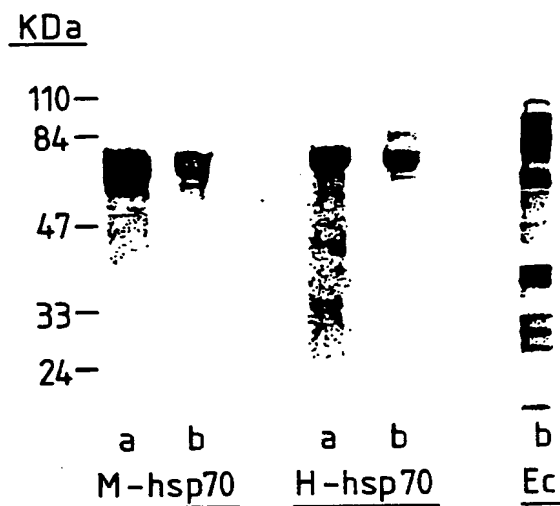


Fig. 3.

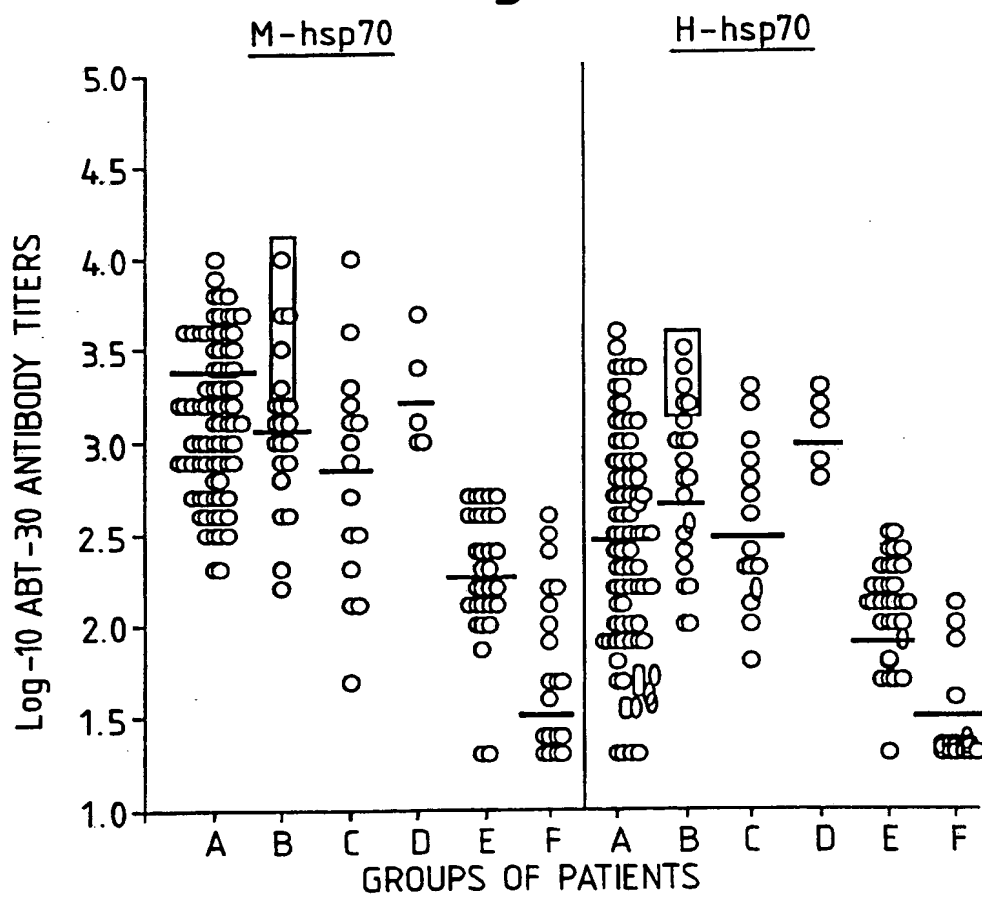


Fig.4.

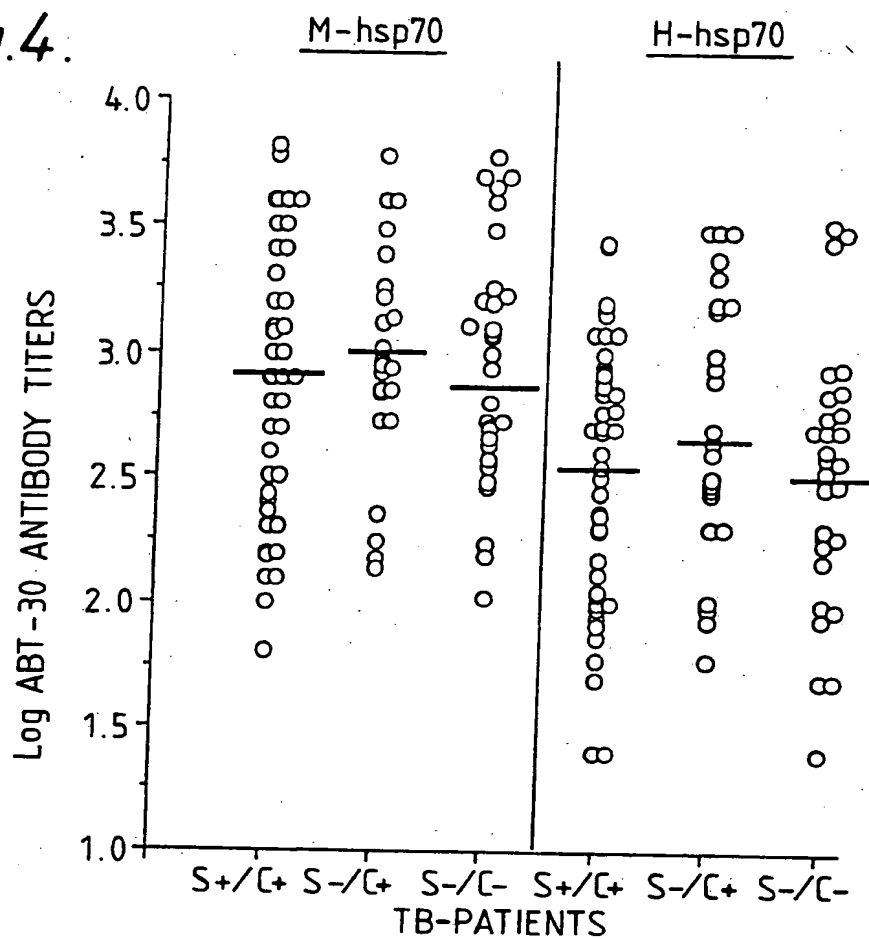


Fig.5.

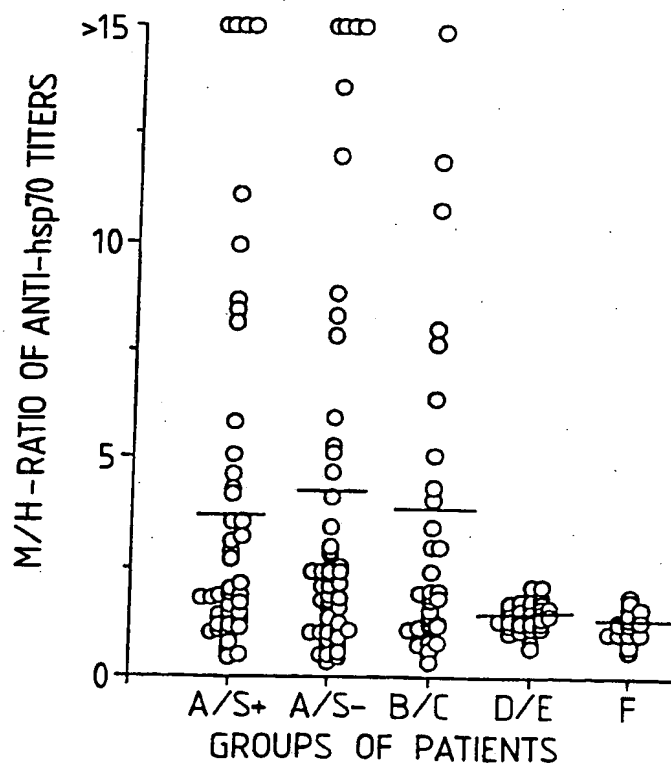


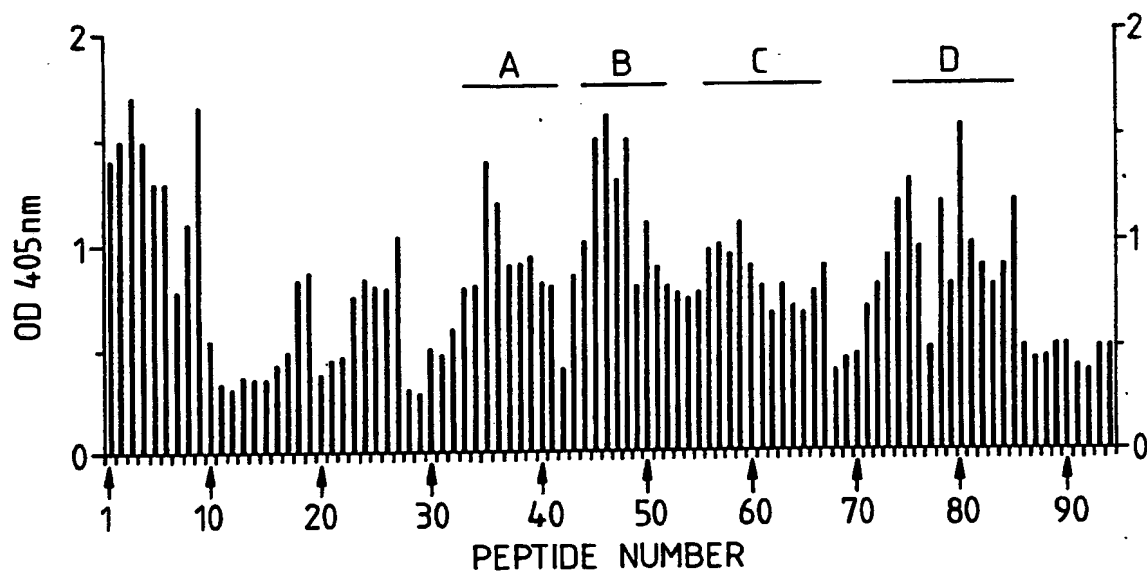
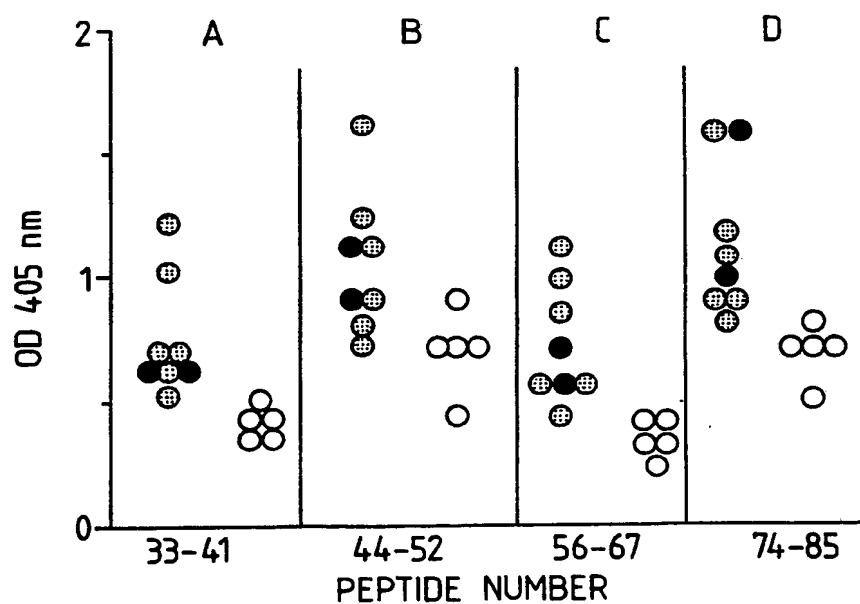
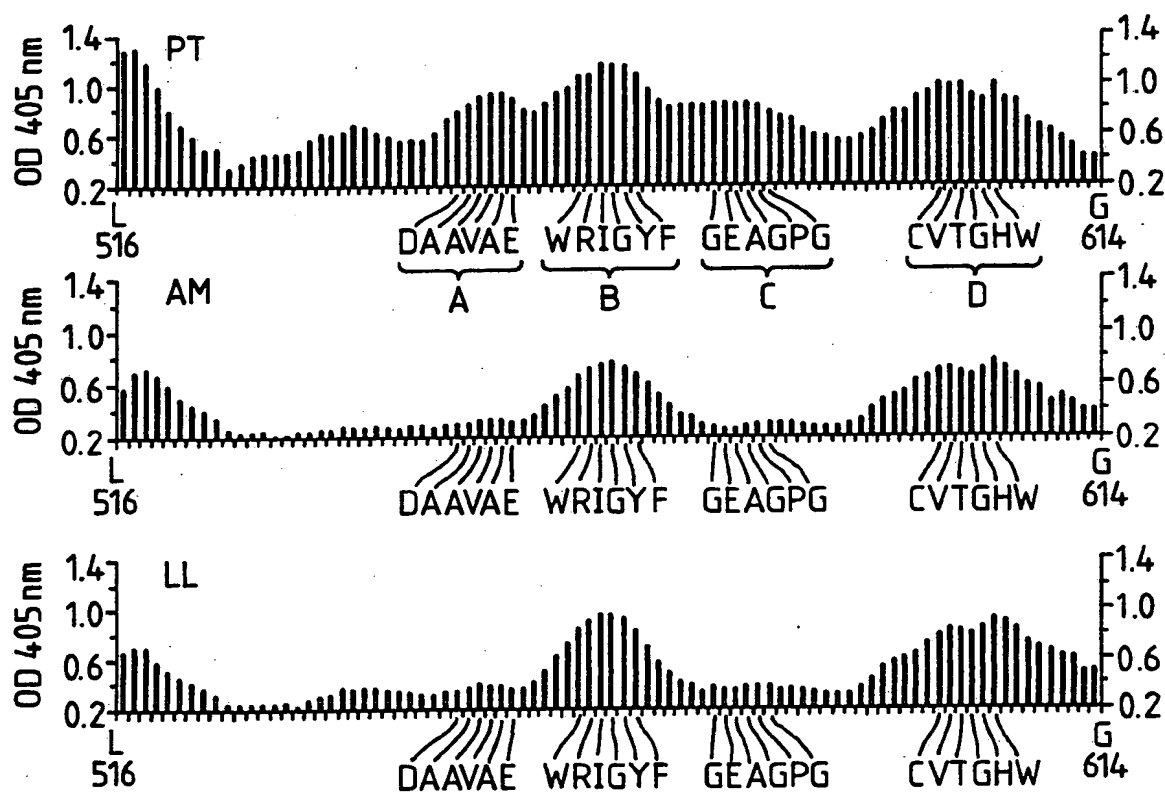
Fig. 6.*Fig. 8.*

Fig. 7.



INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 93/00087

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC Int.Cl. 5 C07K7/06; C07K7/08; C07K7/10; G01N33/569		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
Int.Cl. 5	C07K	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
X	IMMUNOLOGY AND INFECTIOUS DISEASES vol. 1, no. 5, December 1991, pages 323 - 328 A.A.F. ELSAGHIER ET AL. 'ELEVATED ANTIBODY LEVELS TO STRESS PROTEIN HSP70 IN SMEAR-NEGATIVE TUBERCULOSIS' cited in the application see page 324, left column, paragraph 2 - right column, paragraph 2 see page 327, left column, paragraph 2 <div style="text-align: center;">---</div> <div style="text-align: right;">-/--</div>	1-11
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>¹⁰ Special categories of cited documents: ¹⁰</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="width: 45%;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p> </div> </div>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search		Date of Mailing of this International Search Report
11 MAY 1993		1993 05 10
International Searching Authority		Signature of Authorized Officer
EUROPEAN PATENT OFFICE		FUHR C.K.B.

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category *	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.
X	PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA vol. 85, June 1988, WASHINGTON US pages 4267 - 4270 D. YOUNG ET AL. 'Stress proteins are immune targets in leprosy and tuberculosis' see page 4267, right column, last paragraph - page 4268, left column, paragraph 2; figure 1 ---	1-6
P,X	MOLECULAR IMMUNOLOGY vol. 29, no. 9, September 1992, pages 1153 - 1156 A. ELASGHIER ET AL. 'Localisation of linear epitopes at the carboxy-terminal end of the mycobacterial 71 kDA heat shock protein' see the whole document ---	1-11
X	WO,A,8 805 823 (WHITEHEAD INSTITUTE FOR BIOMEDICAL RESEARCH) 11 August 1988 see claims 10,11,13,14,23-27; figure 7 ---	1-6,9-11
A	GB,A,2 239 246 (AKTIEBOLAGET ASTRA) 26 June 1991 see chapter 5.3-5.3.4 on pages 16-19 see claims 1,2,8,13,17,19 -----	1-11

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

GB 9300087
SA 68962

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 11/05/93

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
WO-A-8805823	11-08-88	AU-A-	1548388	24-08-88
		EP-A-	0345299	13-12-89
GB-A-2239246	26-06-91	AU-A-	6807390	27-06-91
		AU-A-	7564491	21-10-91
		CN-A-	1060310	15-04-92
		EP-A-	0479971	15-04-92
		WO-A-	9114448	03-10-91